CENTER FOR DRUG EVALUATION AND RESEARCH

APPLICATION NUMBER: 75706

DRAFT FINAL PRINTED LABELING

GFF 2/21/02

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2003

Drug Facts (continued)

Prug Facts (continued)
 reduces swelling of nasal passages
 temporarily relieves sinus congestion and pressure
 temporarily restores freer breathing through the nose

- temporarily restores freer breathing through the nose
 Warnings
 Do not use
 if you have ever had an allergic reaction to this product or any of its ingredients
 if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking the product.

Drug Facts (continued)

- Ask a doctor before use if you have

 heart disease

 thyroid disease

 trouble urmating due to an enlarged prostate gland

 liver or kidney disease. Your doctor should
 determine if you need a different dose.

When using this product do not take more than directed. Taking more than directed may cause drowsiness.

- Stop use and ask a doctor if

 an allergic reaction to this product occurs. Seek medical help right away.

 symptoms do not improve within 7 days or are accompanied by a fever

 nervousness, dizziness or sleeplessness occurs

Drug Facts (continued)		Drug Facts (continued)
If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children. In case of overdose, get medical help or contact a POSOOC Control Center night units.		Other information safety sealed: sealed with printed foil under cap for your protection. store between 2° and 25°C (36° and 77°F) protect from light and store in a dry place Inactive ingredients black iron oxide, candelilla wax powder, colloidal silicon dioxide, glyceryl monostearate, hydroxypropyl methylcellulose, lactose monohydrate, magnesium stearate, poyethylene glycol, polysor-magnesium stearate, poyethylene glycol, polysor-
Directions do not divide, crush, chew or dissolve the tablet adults and children 12 years and over glass of water; not more than 1 tablet in 24 hours		
children under 12 years of age	ask a doctor	hate 80, propylene glycol, sodium lauryl sulfate, talc and titanium dioxide
consumers with liver or kidney disease	ask a doctor	

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ER Tablets,

10 mg/240 mg

7018 (02/03)

NEX \$2037-605-10

Non-Drowsy*

LORATADINE AND PSEUDOEPHEDRINE SULFATE EXTENDED RELEASE TABLETS (24 Hour Formulation)

Loratadine 10 mg/Antihistamine Pseudoephedrine Sulfate 240 mg/Nasal Decongestant

For Relief of: Nasal and Sinus Congestion Sneezing; Runny Nose Itchy, Watery Eyes Itchy Throat or Nose

Allergy & Congestion

*When taken as directed. See Drug Facts Panel.

1000 EXTENDED RELEASE TABLETS

Manufactured by: Andrx Pharmaceuticals, Inc. Fort Lauderdale, FL 33314

Drug Facts

Active ingredients (in each tablet)

Pseudoephedrine sulfate 240 mg.

Purpose

Uses

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
- runny nose sneezing itchy, watery eyes itching reduces swelling of nasal passages itching of the nose or throat
- temporarily relieves sinus congestion and pressure temporarily restores freer breathing through the nose

Warnings Do not use

- if you have ever had an allergic reaction to this product or any of its ingredients
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking the product.

Ask a doctor before use if you have

- heart disease
- thyroid disease
- high blood pressure
- rouble urinating due to an enlarged prostate gland
- liver or kidney disease. Your doctor should determine if you need a different dose.

When using this product do not take more than directed. Taking more than directed may cause drowsiness.

Stop use and ask a doctor if

 an allergic reaction to this product occurs. Seek medical help right away.

Drug Facts (continued)

- symptoms do not improve within 7 days or are accompanied
- nervousness, dizziness or sleeplessness occurs

If pregnant or breast-feeding, ask a health professional before

Keen out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- do not divide, crush, chew or dissolve the tablet
- adults and children 12 years 1 1 tablet daily with a full glass of water; not more than 1 tablet in

children under 12 years of age ask a doctor

consumers with liver or ask a doctor kidney disease

Other information

- safety sealed: sealed with printed foil under cap for your
- store between 2° and 25°C (36° and 77°F)
 protect from light and store in a dry place

Inactive ingredients

black iron oxide, candelilla wax powder, colloidal silicon dioxide, glyceryl monostearate, hydroxypropyl methylcellulose, lactose monohydrate, magnesium stearate, polyethylene glycol, polysorbate 80, propylene glycol, sodium lauryl sulfate, talc and titanium dioxide



ANDA #75-706 Loratadine & Pseudoephedrine Sulfate ER Tablets, 10 mg/240 mg Final Printed Labeling

Andrew NDC 62037- 605 -01	Drug Facts
Non-Drowsy* LORATADINE AND	Active ingredients (in each tablet) Purpose Loratadine 10 mg
PSEUDOEPHEDRINE SULFATE EXTENDED RELEASE TABLETS (24 Hour Formulation)	Uses temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: nasal congestion runny nose seezing titchy, watery eyes titching of the nose or throat
Loratadine 10 mg/Antihistamine Pseudoephedrine Sulfate 240 mg/Nasal Decongestant	FFR 2 1 2003 7019 (02/0

For Relief of: Nasal and Sinus Congestion Sneezing; Runny Nose Itchy, Watery Eyes Itchy Throat or Nose

Allergy & Congestion

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100 EXTENDED RELEASE TABLETS

When taken as directed.
See Drug Facts Panel
Andrx Pharmaceuticals, Inc.
Fort Lauderdale, FL 33314

7019 (02/03)

THE LABELING